







Upwards



Enhancing the experience

easy-to-use

advanced

and on the rise

Innovation is OUR PASSION but it's your reality THAT DRIVES US.

creos[™] is the Nobel Biocare regenerative product portfolio, built to meet your everyday needs. Join us and experience the carefully designed, easy-to-use, effective solutions.

We aim to be the trusted regenerative partner for you and your patients, because you are what we stand for.

Sideways to your side

effective

creosTM xenoprotect

A membrane with outstanding handling that facilitates bone gain



Outstanding handling^{1,2}

- Does not stick to instruments
- Repositioning in-situ possible
- Low surface expansion when hydrated
- Both sides can face the defect

High mechanical strength^{2,3,4}

- Excellent suture retention
- Highly tear-resistant

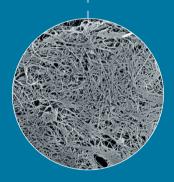
Enduring resistance to degradation in vivo*3

 Manufacturing process intended to preserve the natural structure of the collagen fiber network, to confer a high resistance to degradation

Facilitates bone gain^{2,3,5,6,7,8}

- Excellent tissue compatibility³
- Good clinical results⁵





"What I like is that the handling is very easy. The mechanical stability is very high and when it is rehydrated it adapts very well to the underlying bone"

Dr. Bastian Wessing, Germany



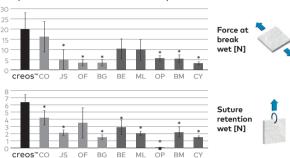
^{*} As shown in an animal model (rat, subcutaneous)

High mechanical strength

In an in vitro study aiming to compare the mechanical strength of commonly used native non-chemically cross-linked and chemically cross-linked collagen membranes4.

- creos[™] xenoprotect demonstrated the highest force at break, wet (21.2 N).
- creos™ xenoprotect had the highest suture retention, wet (6.1 N).

Comparison of commercial membrane in a hydrated state



Non cross-linked collagen membranes (NXL) – CX: creos™ xenoprotect [Nobel Biocare]; CO: Copios [Zimmer]; JS: Jason [botiss]; OF: Osseoguard Flex [3i]; BG: Bio-Gide [Geistlich]

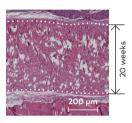
Cross-linked collagen membranes (XL) - BE: BioMend Extend [Zimmer]; ML: Mem-Lok [BioHorizons]; OP: OssixPlus [Datum Dental]; BM: BioMend [Zimmer]; CY: Cytoplast RTM [Osteogenics]

Enduring resistance to degradation in vivo without chemical cross linking³

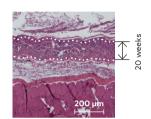
Manufacturing process intended to preserve the natural structure of the collagen fiber network, to confer a high resistance to degradation.3

In an animal model, after 20 weeks, the thickness of xenoprotect decreased only slightly, whereas the reference membrane showed a thickness loss of around 50%, confirming the higher stability of xenoprotect against biodegradation in vivo.3

Representative histological images at 20 weeks implantation in a rat model.







Reference membrane

Facilitates new bone formation^{2,3,5,6,7,8}

New bone formation (%) 34.9% 30 10 Reference

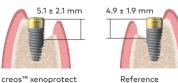
In a comparative in vivo study, creos[™] xenoprotect demonstrated significantly higher new bone formation in the central portion of the defect.

This increase in bone formation was associated with significantly increased expression of the growth factor Bmp2, which has a strong role in osteogenesis.⁷

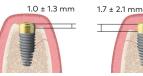
In a randomized controlled clinical trial, 24 patients were treated with creos[™] xenoprotect and 25 with a reference membrane. In the creos[™] xenoprotect group, the defect height reduced at 6-month re-entry by 81%.

In the reference membrane group, the defect height reduced at 6-month re-entry by 62%.5

Schematic showing the defect height prior to treatment and 6 months after GBR



creos[™] xenoprotect



creos™ xenoprotect



Reference



Scan the code for clinical resources.

^{*}Statistically significant

^{*}Statistically significant

creosTM xenogain

3 methods of application to meet all your bone grafting needs

Regenerating bone for 15 years

Three different methods of application:







al le

Syrir

Similar to human bone

- Chemical composition: Ca/P ratio
- Interconnected macropores^{9,10}

Easy handling

- Homogenous particle size⁹
- Hydrophilic for fast rehydration^{11,12}

Solid foundation for dental implant treatment

- Osteoconductive properties¹⁰
- Long-term volume stability¹³
- Uneventful healing^{8,11,12,13,14}



"I appreciated its handling properties and I see its high hydrophilicity as a biological advantage in sinus grafting and peri-implant defect regeneration"

Dr. Werner Zechner, Austria



creos™ xenogain collagen





ck Syrin

Purified cancellous bovine bone mineral granules and 10% porcine collagen in block form and syringe. The collagen helps to hold creos™ xenogain collagen in the desired place. Especially recommended for extraction socket management.





Scaffold for successful regeneration

Preserved natural features of bone through optimized manufacturing process.¹⁰

Chemical composition

With a calcium phosphate ratio that reflects the composition in human bone and a structure with low crystallinity. The body accepts creos™ xenogain as a suitable framework for bone formation.9

Particle size

- Homogenous particle size9
- Maintains space for bone regeneration¹²

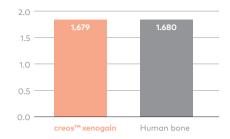
Preserved Nanostructure

Nanostructure preserved thanks to treatment at comparatively low temperature (600°C) and no sintering.¹⁰

Macro and micro-structure

Interconnected macropores allow cells to invade bone grafts and micropores contribute to capillary liquid uptake (hydrophilicity). 15,16

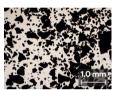
Calcium Phosphate Ratio



Photographic micrograph of creos[™] xenogain and reference product showing the particle size distribution (magnification 20x)







Reference product (0.25 – 1.0 mm)

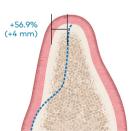
Solid foundation for implant placement

The graft integrates with the newly formed bone, building a basis for successful implant placement.¹²

Schematic showing the defect and bone size prior to and after GBR



Initial situation before GBR

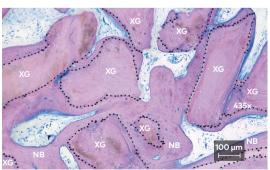


8 months post-surgery

In a multicenter clinical study involving 46 patients, bone increase after 8 months was 4.0 mm (+56.9 % gain) and 4.7 mm (51.0 % gain) at 1 and 3 mm from the top of the crest, respectively.8

GBR led to robust bone regeneration during the 8 months of healing, enabling successful placement of 91 implants in 43 patients, with an average insertion torque of 37.8 ± 5.1 Ncm.8

Histological cross section of the cellular components; NB – new bone, XG – graft, scale bar shown in the bottom right corner, red dashed line: bone to graft particle contact.

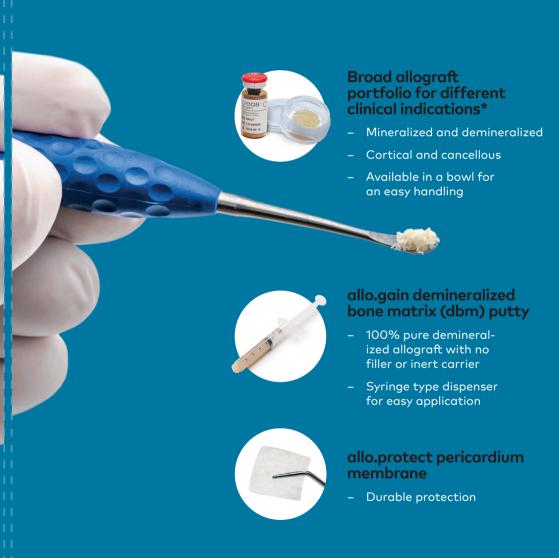


Histological assessment of the trephine cores showed 37.3 % of new bone, 39.1 % of graft material, and 23.6 % of soft tissue (n = 6 cores, 3 patients).8



creos[™] allo.gain & allo.protect

A wide range of allograft materials, because all your cases are different

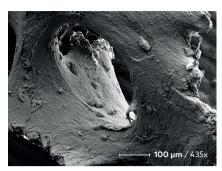


Ensuring safety and quality levels

- A proprietary patented process of tissue cleaning and sterilization
- The tissue bank follows strict processing procedures in order to ensure safe tissue grafts of the highest quality for transplantation



creos™ allo.gain bone particulate: a wide range of options



Mineralized cancellous bone

Mineralized cortical bone

Offers a high density bone with particle size range from 0.125 mm to 1 mm and available volume of 0.25 cc to 2.0 cc.

Mineralized/demineralized cortical bone

Blend of 70% mineralized and 30% demineralized cortical bone.

Mineralized corticocancellous bone

Blend of cortical and cancellous bone produced from sections of the ilium.

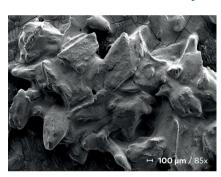
Mineralized cancellous bone

A porous structured bone with particle size range from 0.25 mm to 1 mm and available volume of 0.25 cc to 2.0 cc.

Demineralized cortical bone

Demineralized high density bone with particle size range from 0.125 mm to 1 mm and available volume of 0.25 cc to 2.0 cc.

creos™ allo.gain demineralized bone matrix (dbm) putty



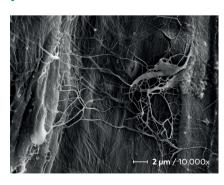
100% pure allograft

The creos[™] allo.gain dbm putty is 100% pure demineralized allograft with no filler or inert carrier.

Available in Three volumes

The creos[™] allo,agin dbm putty is available in three different volumes: 0.5 cc, 1.0 cc and 2.5 cc. This means that the required amount can be used with minimal excess material.

creos™ allo.protect pericardium membrane



Three sizes

creos™ allo.protect is available in three different sizes: 1.0 x 1.0 cm, $1.5 \times 2.0 \text{ cm}$ and $2.0 \times 3.0 \text{ cm}$.

Easy to handle

Easy to tack and suture with high tear resistance. Rapid hydration and easy manipulation. Maintains shape and size once placed.

Durable protection

Strong and stable due to the pore structuwre of native pericardium.

Biocompatible and tissue friendly

Preservation of the native pericardium collagen matrix and its mechanical properties.



Scan the code for clinical resources.

creos[™] syntoprotect

Your easy choice of dense PTFE membranes to expand grafting options



syntoprotect PTFE membrane

Purposely leave the membrane exposed

Preservation of the soft tissue architecture and keratinized mucosa

Non-resorbable

Will not resorb prematurely - you dictate healing time

100% dense (non-expanded) PTFE

Impervious to bacteria - pore size less than 0.3 µm

Soft tissue attaches, but doesn't grow through the membrane

Exposed membrane allows for non-surgical removal; no anesthesia required





Delicate, lightweight framework

Easy to trim and compliant with the overlying soft tissues

Less is more

Н

П

Less titanium bulk allows for greater versatility in shaping and placement, providing additional stability in large, non-spacemaking osseous defects

syntoprotect Ti-reinforced PTFE membrane

Handling options

Broad portfolio with 13 shapes in 2 thicknesses

Traditional frame design

Incorporating delicate and strategically-placed titanium "struts" with more than 25 years of clinical history and successful use in GRB

Unique properties of dense PTFE membranes

Dense PTFE



SEM image courtesy of Schüpbach Ltd, Switzerland.

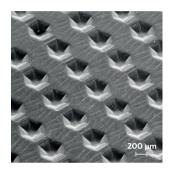
Expanded PTFE



SEM image courtesy of Schüpbach Ltd. Switzerland

Dense PTFE was designed to withstand exposure in the oral environment, which represents an improvement to earlier versions of expanded PTFE in applications such as socket preservation where deliberate membrane exposure offers several advantages.

Designed to aid in membrane stabilization



SEM image courtesy of Schüpbach Ltd, Switzerland.

Hexagonal surface dimples provide a textured surface that increases the area available for cellular attachment without increasing porosity. The textured surface is designed to help stabilize the membrane and the soft tissue flap.



Although PTFE is inherently a non-stick material, cells attach to the outside of the dense PTFE membranes. Cellular adhesion is important to create a seal around the edges of exposed dense PTFE membranes or to support primary closure in larger graft applications.

Clinical evidence

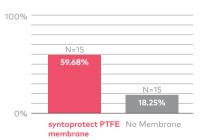
Efficacy

Bone loss 1-year post-extraction¹⁹



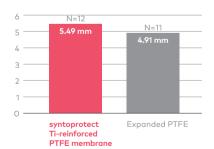
Vertical bone loss measured at crest. Horizontal measured from stent to buccal plate.

Soft tissue regeneration 90 days post-extraction²⁰



Measured as reduction of the occlusal distance between buccal and lingual gingival margins.

Vertical ridge augmentation around implants²¹



Mean vertical bone regeneration.

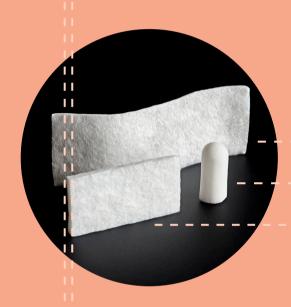
Predictability

In two separate studies treating a total of 696 extraction sites using dense PTFE membranes in an exposed technique, there were no reported infections. 17,18



creosTM absorbable collagen dental wound dressings

Complementing your regenerative set



Three different shapes

creos™ xenotape

creos™ xenoplug

creos™ xenocote



Soft, white, pliable, non-friable, absorbent sponges made from collagen obtained from bovine deep flexor (Achilles) tendons.

Indicated for application to moist or bleeding clean oral wounds created during dental surgery, to control bleeding and protect the surface of the wound from further injury.

Can be used during surgical procedures prior to wound closure or left in-situ.



Depending on the chosen wound dressings product, one or more of the following applications are suitable*:

- Minor oral wounds
- Closure of grafted sites
- Repair of Schneiderian membrane
- Palatal donor sites
- Mucosal flaps
- Extraction sites
- Biopsy sites

Arrive ready to use and are easy to handle.





^{*}See Instructions For Use for full prescribing information, including indications, contraindications, warnings and precautions.

Products

creos[™] xenoprotect

Porcine collagen membrane

Size	Article No.
15x20 mm	N1520
25x30 mm	N2530
30x40 mm	N3040

CreosTM xenogain Deproteinized bovine bone matrix

Weight	Granule size Volume		Vial	Bowl	Syringe
0.25	Small (0.2-1.0 mm)	0.36 cc	N1110	N1110-B	N1210
0.25 g	Large (1.0- 2.0 mm)	0.54 cc	N1111	N1111-B	N1211
0.50 g	Small (0.2-1.0 mm)	0.82 cc	N1120	N1120-B	N1220
	Large (1.0- 2.0 mm)	1.27 cc	N1121	N1121-B	N1221
1.00 g	Small (0.2-1.0 mm)	1.71 cc	N1130	N1130-B	
	Large (1.0- 2.0 mm)	2.69 cc	N1131	N1131-B	
2.00 g	Small (0.2-1.0 mm)	3.64 cc	N1140	N1140-B	
	Large (1.0- 2.0 mm)	5.74 cc	N1141	N1141-B	

creos[™] xenogain collagen creos[™] xenogain + 10% porcine collagen type I

	Size		Article No.
	100 mg	6 x 6 x 6 mm	N1320
Block	250 mg	7 x 8 x 9 mm	N1330
	500 mg	9 x 10 x 11 mm	N1340
Comite and	250 mg	4.6 x 40 mm	N1410
Syringe	500 mg	5.6 x 45 mm	N1420

creos™ allo.gain

bone particulate

Vial	min/demin cortical	miner corticoco		miner cance		mineralized cortical		tical	demineralized cortical	
	Medium 0.25-1 mm	Medium 0.25-1 mm	Large 0.5-1 mm	Medium 0.25-1 mm	Large 0.5-1 mm	Small 0.125-0.85 mm	Medium 0.25-1 mm	Large 0.5-1 mm	Small 0.125-0.85 mm	Large 0.5-1 mm
0.25 cc		N4510	N4511	N4210	N4211	N4110	N4111	N4112	N4310	N4311
0.50 cc	N4410	N4520	N4521	N4220	N4221	N4120	N4121	N4122	N4320	N4321
1.00 cc	N4420	N4530	N4531	N4230	N4231	N4130	N4131	N4132	N4330	N4331
2.00 cc	N4430	N4540	N4541	N4240	N4241	N4140	N4141	N4142	N4340	N4341

Bowl	min/demin cortical	mineralized corticocancellous	mineralized cancellous*	mineralized cortical*
	Medium 0.25-1 mm	Medium 0.25-1 mm	Medium 0.25-1 mm	Medium 0.25-1 mm
0.25 cc		N4510-B	N4210-B	N4111-B
0.50 cc	N4410-B	N4520-B	N4220-B	N4121-B
1.00 cc	N4420-B	N4530-B	N4230-B	N4131-B
2.00 cc	N4430-B	N4540-B	N4240-B	N4141-B

^{*}Available Q2/2022

creos™ allo.gain

dbm putty

Size	Article No.
0.50 cc	N6110
1.00 cc	N6120
2.50 cc	N6130

creosTM allo.protect

Size	Article No.
10 x 10 mm	N7110
15 x 20 mm	N7120
20 x 30 mm	N7140

creos[™] wound dressings

Product type	Configuration/size		Thickness	Article No.
creos™ xenotape	1 in x 3 in	(2.5 cm x 7.5 cm)	0.3-0.8 mm	WD62200
creos™ xenoplug	0.375 in x 0.75 in	(1 cm x 2 cm)		WD62202
creos™ xenocote	0.75 in x 1.5 in	(2 cm x 4 cm)	2-4 mm	WD62201

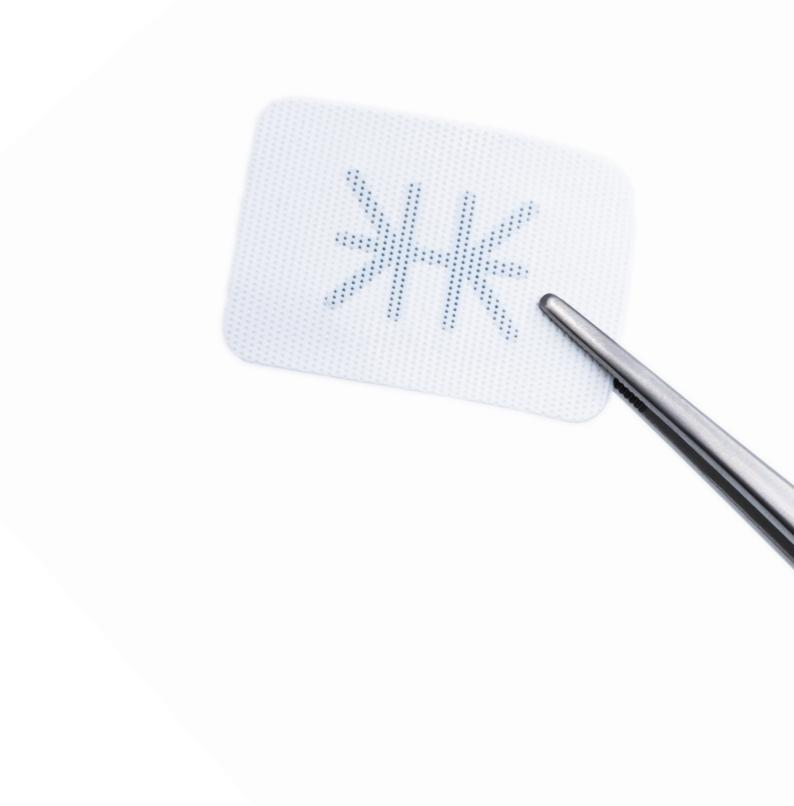
(10 per dispenser)

creos[™] syntoprotect PTFE membrane

Shape	Picture	Size	Thickness	Article No.	Units / box	Description
	25 20	200 μm	N162530-1	1		
Large	Large	25 x 30 mm	200 µm	N162530-4	4	Designed specifically for extraction site grafting and augmentation procedures
C!!	Small 12 x 24 mm	200 µm	N161224-1	1	where exposure to the oral cavity is common	
Small		12 x 24 mm	200 μm	N161224-10	10	

creos[™] syntoprotect Ti-reinforced PTFE membrane

Shape	Picture	Size	Thickness	Article No.	Units / box	Description
NI - 1		12 2/	150 µm	N1615TI-A01-1	1	Designed for narrow single-tooth
No. 1		12 x 24 mm	250 µm	N1625TI-A01-1	1	extraction sites, especially where one bony wall is missing
No. 2		14 x 24 mm	150 µm	N1615TI-A02-1	1	Designed for single-tooth extraction sites, especially where one or
NO. 2		14 X 24 MM	250 µm	N1625TI-A02-1	1	more bony walls are missing
No. 3		17 x 25 mm	150 µm	N1615TI-A03-1	1	Designed for large buccal defects
10. 3		17 X 25 111111	250 µm	N1625TI-A03-1	1	Designed for large boccar defects
lo. 4		20 x 25 mm	150 µm	N1615TI-A04-1	1	Designed for large extraction sites
	7 1	20 X 20 11111	250 µm	N1625TI-A04-1	1	and limited ridge augmentation
lo. 5		36 x 25 mm	150 µm	N1615TI-A05-1	1	Designed for large extraction sites and limited ridge augmentation
NO. 5		30 X 23 IIIIII	250 µm	N1625TI-A05-1	1	in the anterior maxilla
			150 µm	N1615TI-A06-1	1	Designed for large bony defects,
lo. 6		25 x 30 mm	250 µm	N1625TI-A06-1	1	including ridge augmentation
	X		150 µm	N1615TI-A07-1	1	Designed for large bony defects, including
lo. 7	0.7	30 x 41 mm	250 μm	N1625TI-A07-1	1	ridge augmentation in the anterior maxillo
	0.8	30 x 40 mm	150 µm	N1615TI-A08-1	1	Designed for very large bony defects,
NO. 8			250 μm	N1625TI-A08-1	1	including ridge augmentation
1- 0	NV	20 / 0	150 µm	N1615TI-A09-1	1	Designed for very large bony defects,
lo. 9		30 x 40 mm	250 μm	N1625TI-A09-1	1	including ridge augmentation
lo. 10		24 x 38 mm	150 µm	N1615TI-A10-1	1	Designed for large extraction sites,
40. 10		24 X 36 IIIIII	250 µm	N1625TI-A10-1	1	including ridge augmentation
	Н		150 µm	N1615TI-A11-1	1	Designed for large bony defects,
No. 11		38 x 38 mm	250 µm	N1625TI-A11-1	1	including ridge augmentation
	11-11		150 µm	N1615TI-A12-1	1	Designed for large bony defects, including
lo. 12		38 x 38 mm	250 μm	N1625TI-A12-1	1	distal extension of the posterior ridge
	NV		150 µm	N1615TI-A13-1	1	Designed for the largest bony defects,
No. 13		40 x 50 mm	250 μm	N1625TI-A13-1	1	including ridge augmentation



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1-10

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Contact your sales representative or call our customer service team

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